AMENDMENTS TO THE CLAIMS

The following listing of claims replaces all prior versions and listings of claims in the application:

1. (currently amended) An isolated nucleic acid which encodes a phytase having a specific activity of at least about 10 20 U/mg protein,

wherein said specific activity is determined by incubating said phytase in a solution containing about 100 mM maleic acid-Tris-HCl, at a pH of about 5.0 7.5, about 1 mM CaCl₂, and about 1.6 mM sodium phytate at about 37° C. for about 30 minutes,

wherein the isolated nucleic acid hybridizes to SEQ. ID. No. 1 under standard conditions either in 6xSSC, 0.6% SDS, 50° C. overnight for Southern blotting or for PCR: 5 mM Mg²⁺⁺, Taq enzyme, premelting, 94° C. for 2 minutes and 30 cycles of melting at 92° C. for 20 seconds, annealing at 50° C. for 30 seconds and extension at 72° C. for 1 minute.

- 2. (previously presented) The isolated nucleic acid according to claim 1, wherein the nucleic acid is a DNA molecule.
 - 3. (currently amended) A vector comprising:

an isolated DNA molecule which encodes a phytase having a specific activity of at least about $\underline{10}$ 20 U/mg protein,

wherein said specific activity is determined by incubating said phytase in a solution containing about 100 mM maleic acid-Tris-HCl, at a pH of about 5.0 7.5, about 1 mM CaCl₂, and about 1.6 mM sodium phytate at about 37° C. for about 30 minutes,

wherein the isolated DNA molecule hybridizes to SEQ. ID. No. 1 under standard conditions either in 6xSSC, 0.5% SDS, 50° C. overnight for Southern blotting or for PCR: 5 mM Mg²⁻¹, Taq enzyme, premelting, 94° C. for 2 minutes and 30 cycles of melting at 92° C. for 20 seconds, annealing at 50° C. for 30 seconds and extension at 72° C. for 1 minute,

wherein the DNA molecule is functionally linked to regulatory sequences capable of expressing a phytase from said DNA sequence.

- 4. (previously presented) The vector according to claim 3 wherein the, DNA molecule further comprises a leader sequence capable of providing for the secretion of said phytase.
- 5. (currently amended) A <u>An isolated prokaryotic host cell transformed by a nucleic acid</u>, wherein the nucleic acid is an isolated nucleic acid which encodes a phytase having a specific activity of at least about <u>10</u> 20 U/mg protein,

wherein said specific activity is determined by incubating said phytase in a solution containing about 100 mm maleic acid-Tris-HCl, at a pH of about 5.0 7.5, about 1 mM CaCl₂2, and about 1.6 mM sodium phytate at about 37° C. for about 30 minutes,

wherein the isolated nucleic acid hybridizes to SEQ. ID. No. 1 under standard conditions either in 6xSSC, 0.6% SDS, 50° C. overnight for Southern blotting or for PCR: 5 mM Mg²⁺⁺, Taq enzyme, premelting, 94° C. for 2 minutes and 30 cycles of melting at 92° C. for 20 seconds, annealing at 50° C. for 30 seconds and extension at 72° C. for 1 minute.

- 6. (currently amended) A <u>An isolated prokaryotic host cell according to claim 5</u>, wherein the host cell is selected from the group comprising *E. coli*, *Bacillus* sp., *Lactobacillus* sp. and *Lactococcus* sp.
- 7. (currently amended) A An isolated eukaryotic host cell or organism transformed by a nucleic acid, wherein the nucleic acid is an isolated nucleic acid which encodes a phytase having a specific activity of at least about 10 20 U/mg protein,

wherein said specific activity is determined by incubating said phytase in a solution containing about 100 mM <u>maleic acid-Tris-HCl</u>, at a pH of about <u>5.0</u> 7.5, about 1 mM CaCl2, and about 1.6 mM sodium phytate at about 37° C. for about 30 minutes,

wherein the isolated nucleic acid hybridizes to SEQ. ID. No. 1 under standard conditions either in 6xSSC, 0.6% SDS, 50° C. overnight for Southern blotting or for PCR: 5 mM Mg²⁺⁺, Taq enzyme, premelting, 94° C. for 2 minutes and 30 cycles of melting at 92° C. for 20 seconds, annealing at 50° C. for 30 seconds and extension at 72° C. for 1 minute.

- 8. (currently amended) A-An isolated eukaryotic host cell or organism according to claim 7, wherein the host cell is selected from the group comprising *Aspergillus* sp., *Humicola* sp., *Pichia* sp., *Trichoderma* sp. *Saccharomyces* sp. and plants such as soybean, corn and rapeseed.
- 9. (currently amended) A method for the production of phytase comprising: transforming a prokaryotic host cell with an isolated nucleic acid, wherein the isolated nucleic acid encodes a phytase having a specific activity of at least about 10 20 U/mg protein, wherein said specific activity is determined by incubating said phytase in a solution containing about 100 mM maleic acid-Tris-HCl, at a pH of about 5.0 7.5, about 1 mM CaCl₂2, and about 1.6 mM sodium phytate at about 37° C. for about 30 minutes, wherein the isolated nucleic acid hybridizes to SEQ. ID. No. 1 under standard conditions either in 6xSSC, 0.6% SDS, 50°C. overnight for Southern blotting or for PCR: 5 mM·Mg²⁺⁺, Taq enzyme, premelting, 94° C. for 2 minutes and 30 cycles of melting at 92° C. for 20 seconds, annealing at 50° C. for 30 seconds and extension at 72° C. for 1 minute;

culturing or cultivating the prokaryotic host cell under conditions effective for producing phytase; and

recovering phytase.

10. (currently amended) A method for the production identification of a nucleic acid which encodes a phytase, wherein a probe comprising a nucleic acid of SEQ ID NO. 1 or a fragment thereof which encodes a phytase is hybridized to a sample suspected of containing said nucleic acid which encodes a phytase, under standard hydridization conditions either in 6xSSC, 0.6% SDS, 50° C. overnight or functional equivalents thereof for Southern blotting or for PCR 5 mM Mg²⁺⁺, Taq enzyme, premelting, 94° C. for 2 minutes and 30 cycles of metlting at 92° C. for 20 seconds, annealing at 50° C. for 30 seconds and extensioin at 72° C. for 1 minute,

wherein the nucleic acid which encodes a phytase has a specific activity of at least about 10 20 U/mg protein,

wherein said specific activity is determined by incubating said phytase in a solution containing about 100 mM maleic acid-Tris-HCl, at a pH of about 5.0 7.5, about 1 mM CaCl₂2, and about 1.6 mM sodium phytate at about 37° C. for about 30 minutes, wherein the isolated Page 5 of 10

nucleic acid hybridizes to SEQ. ID. No. 1 under standard conditions either in 6xSSC, 0.6% SDS, 50° C. overnight for Southern blotting or for PCR: 5 mM Mg²⁺⁺, Taq enzyme, premelting, 94° C. for 2 minutes and 30 cycles of melting at 92° C. for 20 seconds, annealing at 50° C. for 30 seconds and extension at 72[deg.] C. for 1 minute.

11. (currently amended) A method for the production of phytase comprising: transforming a <u>an isolated</u> eukaryotic host cell with an isolated nucleic acid, wherein the isolated nucleic acid encodes a phytase having a specific activity of at least about 20 U/mg protein, wherein said specific activity is determined by incubating said phytase in a solution containing about 100 mM <u>maleic acid-Tris-HCl</u>, at a pH of about <u>5.0</u> 7.5, about 1 mM CaCl₂, and about 1.6 mM sodium phytate at about 37° C. for about 30 minutes, wherein the isolated nucleic acid hybridizes to SEQ. ID. No. 1 under standard conditions either in 6xSSC, 0.6% SDS, 50° C. overnight for Southern blotting or for PCR: 5 mM Mg²⁺⁺, Taq enzyme, premelting, 94° C. for 2 minutes and 30 cycles of melting at 92° C. for 20 seconds, annealing at 50° C. for 30 seconds and extension at 72° C. for 1 minute;

culturing or cultivating the eukaryotic host cell under conditions effective for producing phytase; and

recovering phytase.

- 12. (new) An isolated nucleic acid of claim 1 wherein the nucleic acid encodes a phytase having a specific activity of at least about 20 U/mg protein and wherein said specific activity is determined by incubating said phytase in a solution containing about 100 mM Tris-HCl at a pH of about 7.5 1 mM CaCl₂, and about 1.6 mM sodium phytate at about 37° C. for about 30 minutes.
- 13. (new) A method for the production of a nucleic acid which encodes a phytase having a specific activity of at least 10 U/mg protein, wherein said specific activity is determined by incubating said phytase in a solution containing about 100 mM maleic acid-Tris, at a pH of about 5.0, about 1 mM CaCl₂, and about 1.6 mM sodium phytate at about 37° C for about 30 minutes, comprising:

providing two or more oligonucleotide primers which hybridize to SEQ ID NO: 1 or to a complement of SEQ ID NO: 1;

contacting the two or more oligonucleotides with a sample suspected of containing a polynucleotide encoding a phytase; and

amplifying the polynucleotide encoding a phytase using the two or more oligonucleotide primers and the polymerase chain reaction, wherein the polymerase chain reaction is carried out under standard conditions: 5mM Mg²⁺⁺, Taq enzyme, premelting, 94° C for 2 minutes, and 30 cycles of melting at 92° C for 20 seconds, annealing at 50° C for 30 seconds, and extension at 72° C for 1 minute.

14. (new) A method for the production of a nucleic acid of claim 13 wherein the nucleic acid encodes a phytase having a specific activity of at least 20 U/mg protein, wherein said specific activity is determined by incubating said phytase in a solution containing about 100 mM Tris-HCl at a pH of about 7.5, 1 mM CaCl₂, and about 1.6 mM sodium phytate at about 37° C. for about 30 minutes.